Reference number(s)
3029-A

SPECIALTY GUIDELINE MANAGEMENT

DUOPA (carbidopa and levodopa enteral suspension)

POLICY

I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

FDA-Approved Indication
Duopa is indicated for the treatment of motor fluctuations in patients with advanced Parkinson's disease.

All other indications are considered experimental/investigational and not medically necessary.

II. CRITERIA FOR INITIAL APPROVAL

Parkinson's disease
Authorization of 6 months may be granted for treatment of motor fluctuations in members with advanced Parkinson's disease when all of the following criteria are met:
A. Member is levodopa responsive with clearly defined "on" periods
B. The member has "off" periods greater than 3 hours per day despite optimization efforts
C. The member must have had an inadequate response or intolerable adverse event with oral carbidopa-levodopa and one of the following anti-Parkinson agents:
    1. Dopamine agonists (e.g., pramipexole, ropinirole)
    2. Monoamine oxidase B (MAO)-B inhibitor (e.g., selegiline, rasagiline)
    3. Catechol-O-methyl transferase (COMT) inhibitor (e.g., entacapone, tolcapone)

III. CONTINUATION OF THERAPY

Authorization of 12 months may be granted for continued treatment in members requesting reauthorization for advanced Parkinson's disease who have demonstrated a positive clinical response to Duopa therapy.

IV. REFERENCES

1. Duopa [package insert]. North Chicago, IL: AbbVie, Inc; March 2022.
2. C. Warren Olanow, Karl Keiburtz, Per Odin, et al. Double blind, double dummy, randomized study of continuous intrajejunal infusion of levodopa-carbidopa intestinal gel in advanced Parkinson's disease, *Lancet Neurol*. 2014 February; 13 (2): 141-149.